

PATENT COOPERATION TREATY

From the
INTERNATIONAL SEARCHING AUTHORITY

To:
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PCT

WRITTEN OPINION OF THE
INTERNATIONAL SEARCHING AUTHORITY

(PCT Rule 43bis.1)

		Date of mailing (day/month/year)	06 NOV 2007
Applicant's or agent's file reference 19025.021		FOR FURTHER ACTION See paragraph 2 below	
International application No. PCT/US04/20751	International filing date (day/month/year) 28 June 2004 (28.06.2004)	Priority date (day/month/year) 24 May 2004 (24.05.2004)	
International Patent Classification (IPC) or both national classification and IPC IPC(8): C12Q 1/68(2006.01);C07H 21/04 USPC: 435/6,69.1,320.1,325;530/350;536/23.5			
Applicant PTC THERAPEUTICS			

1. This opinion contains indications relating to the following items:

- Box No. I Basis of the opinion
- Box No. II Priority
- Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- Box No. IV Lack of unity of invention
- Box No. V Reasoned statement under Rule 43bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- Box No. VI Certain documents cited
- Box No. VII Certain defects in the international application
- Box No. VIII Certain observations on the international application

2. **FURTHER ACTION**

If a demand for international preliminary examination is made, this opinion will be considered to be a written opinion of the International Preliminary Examining Authority ("IPEA") except that this does not apply where the applicant chooses an Authority other than this one to be the IPEA and the chosen IPEA has notified the International Bureau under Rule 66.1bis(b) that written opinions of this International Searching Authority will not be so considered.

If this opinion is, as provided above, considered to be a written opinion of the IPEA, the applicant is invited to submit to the IPEA a written reply together, where appropriate, with amendments, before the expiration of 3 months from the date of mailing of Form PCT/ISA/220 or before the expiration of 22 months from the priority date, whichever expires later.

For further options, see Form PCT/ISA/220.

3. For further details, see notes to Form PCT/ISA/220.

Name and mailing address of the ISA/ US Mail Stop PCT, Attn: ISA/US Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450 Facsimile No. (571) 273-3201	Date of completion of this opinion 15 October 2007 (15.10.2007)	Authorized officer Stephanie K. Mummert, Ph.D. Telephone No. 571-272-0872
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Box No. I Basis of this opinion

1. With regard to the **language**, this opinion has been established on the basis of:
 the international application in the language in which it was filed
 a translation of the international application into _____, which is the language of a translation furnished for the purposes of international search (Rules 12.3(a) and 23.1(b)).
2. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application and necessary to the claimed invention, this opinion has been established on the basis of:
 - a. type of material
 a sequence listing
 table(s) related to the sequence listing
 - b. format of material
 on paper
 in electronic form
 - c. time of filing/furnishing
 contained in the international application as filed.
 filed together with the international application in electronic form.
 furnished subsequently to this Authority for the purposes of search.
3. In addition, in the case that more than one version or copy of a sequence listing and/or table(s) relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.
4. Additional comments:

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Box No. IV Lack of unity of invention

1. In response to the invitation (Form PCT/ISA/206) to pay additional fees the applicant has, within the applicable time limit:
 paid additional fees
 paid additional fees under protest and, where applicable, the protest fee
 paid additional fees under protest but the applicable protest fee was not paid
 not paid additional fees
2. This Authority found that the requirement of unity of invention is not complied with and chose not to invite the applicant to pay additional fees.
3. This Authority considers that the requirement of unity of invention in accordance with Rule 13.1, 13.2 and 13.3 is
 complied with
 not complied with for the following reasons:

See the lack of unity section of the International Search Report (Form PCT/ISA/210)

4. Consequently, this opinion has been established in respect of the following parts of the international application:

all parts.
 the parts relating to claims Nos. 1-27

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Box No. V Reasoned statement under Rule 43 bis.1(a)(i) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N) Claims NONE YES
 Claims 1-27 NO

Inventive step (IS) Claims NONE YES
 Claims 1-27 NO

Industrial applicability (IA) Claims 1-27 YES
 Claims NONE NO

2. Citations and explanations:

Please See Continuation Sheet

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Supplemental Box

In case the space in any of the preceding boxes is not sufficient.

V. 2. Citations and Explanations:

Claim Interpretation

The term 'in an absence of SEQ ID NO:4' is being given the broadest reasonable interpretation in light of the specification. The term is not explicitly defined in the spec. Instead, SEQ ID NO:4 is referred to as NeRP and as "SEQ ID NO:4 sets forth a NeRP1, a 336 nucleotide region of a VEGF 5'UTR" (p. 8 of specification) and it is noted that searching the sequence against nucleotide databases does not necessarily provide art where the sequence is deleted. However, the nucleotide boundaries of SEQ ID NO:4 are not established relative to the context of the overall full-length VEGF 5' UTR. Therefore, without clear nucleotide boundaries of the region comprising SEQ ID NO:4, the term is being interpreted as reading on art where the 5' UTR is deleted partially, either at the 5' end of the UTR, the 3' end of the UTR or from the middle.

The term 'UTR having a NeRP1 (SEQ ID NO: 4)' is also being given the broadest reasonable interpretation in light of the specification. As noted above, the limitations of SEQ ID NO:4 are not clearly defined. The term is being interpreted as the opposite of 'in the absence of SEQ ID NO:4' and is interpreted as reading on art where a full length VEGF 5' UTR is present in the nucleotide construct.

The limitations of SEQ ID NO:3 are also not explicitly defined in the spec. Instead, SEQ ID NO:3 is referred to as PTCRE1 and as "SEQ ID NO:3 sets forth a PTCRE1, a 702 nucleotide region of VEGF 5'UTR" (p. 8 of specification) and like SEQ ID NO:4, the nucleotide boundaries of SEQ ID NO:3 are not established relative to the full-length 5' UTR and searching the sequence against nucleotide databases does not necessarily provide art where the sequence is deleted. Therefore, the term 'wherein the PTCRE is not SEQ ID NO:3' is being interpreted as reading on art the term is being interpreted as reading on art where the 5' UTR is partially deleted, either at the 5' end of the UTR, the 3' end of the UTR or from the middle. And where SEQ ID NO:3 is not described either way, or particularly where the sequence comprising 'SEQ ID NO:3, a fragment thereof, or a complement of either' is being interpreted as reading on art where the full length 5' UTR is present in the nucleotide construct.

Claims 1-27 lack novelty under PCT Article 33(2) as being anticipated by Forsythe et al. (*Molecular and Cellular Biology*, 1996, vol. 16, no. 9, p. 4604-4613). Forsythe teaches a method of analyzing the effect of hypoxia inducible factor on the expression of VEGF (Abstract).

With regard to claims 1-10 and 14, Forsythe teaches a variety of nucleic acid constructs and nucleic acids that comprise a nucleic acid encoding a reporter polypeptide, wherein the nucleic acid sequence encoding a reporter polypeptide is operably linked to a NeRP, said NeRP (SEQ ID NO:4) is operably linked to a PTCRE (wherein said PTCRE is not SEQ ID NO:3), and expression of said reporter polypeptide is capable of being modulated relative to in an absence of said NeRP (Figure 1A, where a variety of constructs comprising deletions of the 5' UTR, and therefore in the absence of SEQ ID NO:3; see also p. 4605, col. 1, 'reporter plasmid constructs')

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In case the space in any of the preceding boxes is not sufficient.

heading, where the UTR is linked to luciferase reporter gene).

With regard to claims 11-13 and 15-21, Forsythe teaches a reporter construct wherein said VEGF 5' UTR is in an absence of SEQ ID NO:4 and contains an intron (Figure 1A, where a variety of constructs comprising deletions of the 5' UTR, and therefore in the absence of SEQ ID NO:4; see also p. 4605, col. 1, 'reporter plasmid constructs' heading, where the UTR is linked either to VEGF ORF or luciferase reporter gene), wherein these constructs produce polypeptides (see also p. 4605, col. 1, 'reporter plasmid constructs' heading, where the UTR is linked either to VEGF ORF or luciferase reporter gene), and are produced *in vitro* (p. 4605, where the constructs are produced *in vitro*, see 'transient expression assays' heading).

With regard to claims 22-24, Forsythe teaches a nucleic acid molecule that comprises 95-99% sequence identity with a nucleic acid molecule of SEQ ID NO:3, a fragment thereof or a complement of either, consists of SEQ ID NO:3 or a fragment or complement thereof, or consists of a nucleic acid linked to a reporter polypeptide wherein the nucleic acid sequence consists of SEQ ID NO:3 (Figure 1A, where a variety of constructs comprising deletions and full length versions, see KpnI of the 5' UTR, and therefore comprising SEQ ID NO:3).

With regard to claims 23-27, Forsythe teaches a nucleic acid molecule that comprises 95-99% sequence identity with a nucleic acid molecule of SEQ ID NO:4, a fragment thereof or a complement of either, consists of SEQ ID NO:4 or a fragment or complement thereof, or consists of a nucleic acid linked to a reporter polypeptide wherein the nucleic acid sequence consists of SEQ ID NO:4 (Figure 1A, where a variety of constructs comprising deletions and full length versions, see KpnI of the 5' UTR, and therefore comprising SEQ ID NO:4).

Claims 22-27 lack novelty under PCT Article 33(2) as being anticipated by Kamiya et al. (US Patent 6,057,437; May 2000) teach the specific nucleotide sequences of VEGF 3' and 5' UTR regions (Table I, col. 10).

With regard to claims 22-24, Kamiya teaches a nucleic acid molecule that comprises 95-99% sequence identity with a nucleic acid molecule of SEQ ID NO:3, a fragment thereof or a complement of either, consists of SEQ ID NO:3 or a fragment or complement thereof, or consists of a nucleic acid linked to a reporter polypeptide wherein the nucleic acid sequence consists of SEQ ID NO:3 (Table 1, col. 10, see sequence alignment below).

Qy	1	TCCAGAGAGAACGAGGGAAAGAGAGAGACGGGGTCAGAGAGAGCGCGCGGGGTGCGAGC	60
Db	337	TCCAGAGAGAACGAGGGAAAGAGAGAGACGGGGTCAGAGAGAGCGCGCGGGGTGCGAGC	396
Qy	61	AGCGAAAGCGACAGGGCAAAGTGAGTGACCTGCTTTGGGGGTGACCGCGGGAGCGCGG	120
Db	397	AGCGAAAGCGACAGGGCAAAGTGAGTGACCTGCTTTGGGGGTGACCGCGGGAGCGCGG	456
Qy	121	CGTCAGCCCTCCCCCTGGATCCCGCAGCTGACCAGTCGCCTGACGGACAGACAGACA	180
Db	457	CGTGAAGCCCTCCCCCTGGATCCCGCAGCTGACCAGTCGCCTGACGGACAGACAGACA	516
Qy	181	GACACCGCCCCCAGCCCCAGCTACCACCTCTCCCCGGCCGGCGGGACACTGGACCG	240
Db	517	GACACCGCCCCCAGCCCCAGCTACCACCTCTCCCCGGCCGGCGGGACACTGGACCG	576
Qy	241	GCGGCCAGCCGGGGAGGGCCGGAGCCCGCGCCGGAGGGGGCTGGAGGGGGTCGGG	300
Db	577	GCGGCCAGCCGGGGAGGGCCGGAGCCCGCGCCGGAGGGGGCTGGAGGGGGTCGGG	636
Qy	301	GCTCGCGCGTCGCAGTGAAACTTTCTGCTCAACTTCTGGCTGTTCTCGCTTCGGAGGA	360
Db	637	GCTCGCGCGTCGCAGTGAAACTTTCTGCTCAACTTCTGGCTGTTCTCGCTTCGGAGGA	696
Qy	361	GCCGTGGTCCCGCGGGGGAGCCGAGCCGAGCGGAGGGAGAAGTGCCTAGCTCGGGC	420
Db	697	GCCGTGGTCCCGCGGGGGAGCCGAGCCGAGCGGAGGGAGAAGTGCCTAGCTCGGGC	756
Qy	421	CGGGAGGAGCCGAGCCCGAGGAGGGAGGGAGGGAGAAGAAGAGAAGGGAGAGGGGG	480
Db	757	CGGGAGGAGCCGAGCCCGAGGAGGGAGGGAGGGAGAAGAAGAGAAGGGAGAGGGGG	816
Qy	481	CCGCAGTGGCAGCTGGCGCTCGGAAGGCCGGCTCATGGACGGGTGAGGCCGGGTGTC	540
Db	817	CCGCAGTGGCAGCTGGCGCTCGGAAGGCCGGCTCATGGACGGGTGAGGCCGGGTGTC	876
Qy	541	GCAGACAGTGTCCAGCCGCGCGCTCCCCAGGCCCTGGCCCGGGCTCGGCCGGGGA	600
Db	877	GCAGACAGTGTCCAGCCGCGCGCTCCCCAGGCCCTGGCCCGGGCTCGGCCGGGGA	936
Qy	601	GGAAGAGTAGCTCGCCGAGGCCCGAGGAGAGCGGGCCGCCACAGCCCGAGCCGGAGA	660
Db	937	GGAAGAGTAGCTCGCCGAGGCCCGAGGAGAGCGGGCCGCCACAGCCCGAGCCGGAGA	996

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QY	661 GGGAGCGCGAGCCGCGCCGGCCCCGGTCGGGCCTCGAAACC 702
Db	997 GGGAGCGCGAGCCGCGCCGGCCCCGGTCGGGCCTCGAAACC 1038

With regard to claims 23-27, Kamiya teaches a nucleic acid molecule that comprises 95-99% sequence identity with a nucleic acid molecule of SEQ ID NO:4, a fragment thereof or a complement of either, consists of SEQ ID NO:4 or a fragment or complement thereof, or consists of a nucleic acid linked to a reporter polypeptide wherein the nucleic acid sequence consists of SEQ ID NO:4 (Table 1, col. 10, see sequence alignment below).

QY	1 TCGCCGGAGGCTTGGGGCAGCCGGTAGCTCGGAGGTCTGGCGCTGGGGGCTAGCACCAAG 60
Db	1 TCGCCGGAGGCTTGGGGCAGCCGGTAGCTCGGAGGTCTGGCGCTGGGGGCTAGCACCAAG 60
QY	61 CGCTCTGCGGGAGGCCAGCGGTTAGGTGGACCGGTCAAGCGGACTCACCGGCCAGGGCG 120
Db	61 CGCTCTGCGGGAGGCCAGCGGTTAGGTGGACCGGTCAAGCGGACTCACCGGCCAGGGCG 120
QY	121 CTCGGTGCTGGAATTGATATTCAATTGATCCGGTTTATCCCTCTTCTTTCTTAAA 180
Db	121 CTCGGTGCTGGAATTGATATTCAATTGATCCGGTTTATCCCTCTTCTTTCTTAAA 180
QY	181 CATTTTTTTAAACTGTATTGTTCTCGTTTAATTATTTTGTGCTGCCATTCCCCA 240
Db	181 CATTTTTTTAAACTGTATTGTTCTCGTTTAATTATTTTGTGCTGCCATTCCCCA 240
QY	241 CTTGAATCGGGCCGACGGCTTGGGGAGATTGCTCTACTTCCCAAATCACTGTGGATTT 300
Db	241 CTTGAATCGGGCCGACGGCTTGGGGAGATTGCTCTACTTCCCAAATCACTGTGGATTT 300
QY	301 GGAAACCAGCAGAAAGAGGAAAGACGCTAGCAAGAGC 336
Db	301 GGAAACCAGCAGAAAGAGGAAAGAGCTAGCAAGAGC 336

Claims 1-27 meet the criteria set out in PCT Article 33(4), and thus have industrial applicability because the subject matter claimed can be made or used in industry.